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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/517,320	07/21/2005	Per Mansson	Mans3011/REF	3650
23364 BACON & TH	7590 01/24/2007 OMAS, PLLC		EXAMINER	
625 SLATERS	LANE		JUNG, UNSU	
FOURTH FLOOR ALEXANDRIA, VA 22314			ART UNIT	PAPER NUMBER
	,		1641	
SHORTENED STATUTOR	Y PERIOD OF RESPONSE	MAIL DATE	DELIVERY MODE	
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Please find below and/or attached an Office communication concerning this application or proceeding.

If NO period for reply is specified above, the maximum statutory period will apply and will expire 6 MONTHS from the mailing date of this communication.

	Application No.	Applicant(s)				
Office Action Comment	10/517,320	MANSSON ET AL.				
Office Action Summary	Examiner	Art Unit				
	Unsu Jung	1641				
The MAILING DATE of this communication appears on the cover sheet with the correspondence address Period for Reply						
A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 3 MONTH(S) OR THIRTY (30) DAYS, WHICHEVER IS LONGER, FROM THE MAILING DATE OF THIS COMMUNICATION. - Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication. - If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication. - Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133). Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).						
Status	•	·				
1) Responsive to communication(s) filed on 30 Oc	ctober 2006.					
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closed in accordance with the practice under E	·					
Disposition of Claims	,	,				
4)⊠ Claim(s) <u>1-12</u> is/are pending in the application.						
4a) Of the above claim(s) <u>8-12</u> is/are withdrawn from consideration.						
5) Claim(s) is/are allowed.						
6)⊠ Claim(s) <u>1-7</u> is/are rejected.						
7) Claim(s) is/are objected to.						
,= ,,	4					
Application Papers	•					
9) The specification is objected to by the Examiner.						
10) \boxtimes The drawing(s) filed on <u>30 October 2006</u> is/are: a) \boxtimes accepted or b) \square objected to by the Examiner.						
Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).						
Replacement drawing sheet(s) including the correction is required if the drawing(s) is objected to. See 37 CFR 1.121(d).						
11) The oath or declaration is objected to by the Examiner. Note the attached Office Action or form PTO-152.						
Priority under 35 U.S.C. § 119	·	•				
12)⊠ Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f). a)⊠ All b)□ Some * c)□ None of:						
1. Certified copies of the priority documents have been received.						
2. Certified copies of the priority documents have been received in Application No						
3. Copies of the certified copies of the priority documents have been received in this National Stage						
application from the International Bureau (PCT Rule 17.2(a)).						
* See the attached detailed Office action for a list of the certified copies not received.						
·		•				
Attachment(s)	•					
1) Notice of References Cited (PTO-892)	4) Interview Summary	(PTO-413)				
2) D Notice of Draftsperson's Patent Drawing Review (PTO-948)	Paper No(s)/Mail Da	ite				
3) Information Disclosure Statement(s) (PTO/SB/08) Paper No(s)/Mail Date 10/30/06.	5) Notice of Informal P 6) Other:	atent Application				
1 3201 140(3)/Wall Date 10/30/00.	J/ [

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DETAILED ACTION

Response to Amendment

- 1. Applicants' amendments to the specification and drawings in the reply filed on October 30, 2006 has been acknowledged and entered.
- 2. Applicants' amendment to claims 1 and 3-5 in the reply filed on October 30, 2006 has been acknowledged and entered.
- 3. Claims 1-12 are pending and claims 1-7 are under consideration for their merits.

Objections Withdrawn

- 4. Applicants' arguments, see p8, filed on October 30, 20006, with respect to the objection of the drawings have been fully considered and are persuasive. The objection of the drawings has been withdrawn in light of the amended Fig. 9 in the reply filed on October 30, 2006.
- 5. Applicants' arguments, see pp8-9, filed on October 30, 20006, with respect to the objection of the specification have been fully considered and are persuasive. The objection of the specification has been withdrawn in light of the specification in the reply filed on October 30, 2006.

Information Disclosure Statement

6. The information disclosure statement (IDS) submitted on October 30, 2006 has been considered by the examiner.

Claim Rejections - 35 USC § 112

- 7. The following is a quotation of the second paragraph of 35 U.S.C. 112:
 - The specification shall conclude with one or more claims particularly pointing out and distinctly claiming the subject matter which the applicant regards as his invention.
- 8. Claims 1-7 are rejected under 35 U.S.C. 112, second paragraph, as being indefinite for failing to particularly point out and distinctly claim the subject matter which applicant regards as the invention.
- 9. In claim 1, the term "amide-group" in line 4 is vague and indefinite. It is unclear whether or not the term "amide-group" in line 4 is referring to "an amide group" in line 2. For the purpose of examination, the term "amide-group" in line 4 has been interpreted as being different from "an amide group" in line 2.

Applicants' arguments, see pp9, filed on October 30, 20006, with respect to the rejection under 35 U.S.C. 112, second paragraph (items 9 and 10 in the Office Action dated June 28, 2006) have been fully considered and are persuasive. However, the term "amide-group" in line 4 of claim 1 remains vague and indefinite as discussed above. Therefore the rejection of claims 1-7 under 35 U.S.C. 112, second paragraph has been maintained.

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Claim Rejections - 35 USC § 103

- 10. The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:
 - (a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negatived by the manner in which the invention was made.
- 11. The factual inquiries set forth in *Graham* v. *John Deere Co.*, 383 U.S. 1, 148 USPQ 459 (1966), that are applied for establishing a background for determining obviousness under 35 U.S.C. 103(a) are summarized as follows:
 - 1. Determining the scope and contents of the prior art.
 - 2. Ascertaining the differences between the prior art and the claims at issue.
 - 3. Resolving the level of ordinary skill in the pertinent art.
 - 4. Considering objective evidence present in the application indicating obviousness or nonobviousness.
- 12. This application currently names joint inventors. In considering patentability of the claims under 35 U.S.C. 103(a), the examiner presumes that the subject matter of the various claims was commonly owned at the time any inventions covered therein were made absent any evidence to the contrary. Applicant is advised of the obligation under 37 CFR 1.56 to point out the inventor and invention dates of each claim that was not commonly owned at the time a later invention was made in order for the examiner to consider the applicability of 35 U.S.C. 103(c) and potential 35 U.S.C. 102(e), (f) or (g) prior art under 35 U.S.C. 103(a).

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13. Claims 1, 2, 6, and 7 are rejected under 35 U.S.C. 103(a) as being unpatentable over Willner et al. (WO 00/43774, July 27, 2000) in view of Svedhem et al. (*J. Org. Chem.* 2001, Vol. 66, pp4494-4503) and Bentley et al. (U.S. PG Pub. No. US 2001/0027212 A1, Oct. 4, 2001).

Willner et al. teaches a sensitive method of detecting small amount of low molecular weight compounds (typically below about 1,500 Daltons), which includes explosive molecules such as DNT and TNT (derivatized explosives, Fig's 1A and 1B) and drugs such as heroin and cocaine (p5, lines 1-15) using quartz crystal microbalance (QCM). Willner et al. further teaches that any method intended for sensing the presence of explosive molecules or other types of low molecular weight molecules such as drugs should be highly sensitive and adapted for detecting a small amount of molecules. The QCM includes a piezoelectric crystal sandwiched between two gold electrodes (Abstract and p23, lines 14-17) coated with an antigen, which is then contacted with an antibody (p24, lines 8-12). Measurement of resonance frequency at this stage yields a certain basic frequency (p24, lines 9-12). Challenging the electrode with a sample comprising antigens causes release of some of the antibodies to yield a soluble antigen-antibody complex, which reduces the immobilized mass and consequently the frequency is increased as a result of and signifies the presence of the assayed molecule in the medium (p24, lines 13-19). However, Willner et al. fails to teach a coated metal surface further comprising a self-assembled monolayer (SAM) of oligo(ethylene glycol)-terminated (OEG-terminated) alkanethiol amides.

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Svedhem et al. teaches a self-assembled monolayer (SAM) of oligo(ethylene glycol)-terminated (OEG-terminated) alkanethiol amides on gold coated surface on a solid support (p4503, right column, Preparation of SAMs) designed to address structure and stability of biosensing interfaces (Abstract). SAM-forming OEG molecules includes alkyl portion of the alkanethiols having 2, 5, 11, and 15 CH₂ groups (methylene groups) and OEG portion has 1, 2, 4, 6, 8, 10, and 12 (CH₂CH₂O) (ethylene oxy) units (Abstract). Organic modifications of gold surfaces by SAMs have proven to be successful in biosensor applications (p4494, *Introduction*, second paragraph). Furthermore, ethylene glycols provide good anchors for biological receptors and ligands and reduce nonspecific binding of proteins and other bioactive molecules (p4494, *Introduction*, first paragraph). Poly(ethylene glycol) derivatives are also ideal as spacer candidates because they are inexpensive, water soluble, stable, and available in a wide range of molecular weight distributions (p4494, *Introduction*, first paragraph). However, Svedhem et al. fails to teach low molecular weight antigens bound via an amide group to the SAM-forming OEG molecules.

Bentley et al. teaches that conventional amide linkages formed between amine groups on drugs, which include peptides, proteins and small agents (antigens), having amine groups and PEG through non-hydrolyzable amide linkages, which are generally stable (p1, paragraph [0007]). However, Bentley et al. fails to teach that antigens are reversibly bound to antibodies specific for the antigens.

Therefore, it would have been obvious to one of ordinary skill in the art at the time of the invention to employ SAM of OEG-terminated alkanethiol amides of Svedhem

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et al. in the QCM biosensor of Willner et al. in order to provide a biosensing interface with structurally stable SAM, which reduce nonspecific binding of proteins and other bioactive molecules. The advantage of having a structurally stable SAM, which has the characteristic of reducing nonspecific binding of proteins and other bioactive molecules provides the motivation to include the SAM of OEG-terminated alkanethiol amides of Svedhem et al. in the QCM biosensor of Willner et al. with a reasonable expectation of success since the solid support of Willner et al. includes a gold coated surface and Svedhem et al. teaches that he SAM of OEG-terminated alkanethiol amides can be formed on gold coated surfaces for use as a biosensing interfaces. Furthermore, it would have been obvious to one of ordinary skill in the art at the time of the invention to use conventional amide linkages formed between amine groups on drugs and ethylene glycol of OEG as taught by Bentley et al. in order to immobilize antigens of interest on the SAM of OEG-terminated alkanethiol amides of Svedhem et al. as the amide linkages are generally stable and non-hydrolyzable. The advantage of amide linkages, which are stable and non-hydrolyzable provides the motivation to employ amide linkages to immobilize antigens of Willner et al. on the SAM of OEG-terminated alkanethiol amides of Svedhem et al. with a reasonable expectation of success as Bentley et al. teaches that small molecules such as drugs can be immobilized to ethylene glycols of PEG, which are also present in OEGs.

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14. Claims 3-5 are rejected under 35 U.S.C. 103(a) as being unpatentable over Willner et al. (WO 00/43774, July 27, 2000) in view of Svedhem et al. (*J. Org. Chem.*

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2001, Vol. 66, pp4494-4503) and Bentley et al. (U.S. PG Pub. No. US 2001/0027212 A1, Oct. 4, 2001) as applied to claim 1 above, and further in view of Duffy (U.S. PG Pub. No. US 2002/0028463 A1, Mar. 7, 2002).

Willner et al. in view of Svedhem et al. and Bentley et al. teaches a coated metal surface on a solid support as discussed above. Willner et al. further teaches that antigens are selected from a group consisting of explosives and narcotics (p5, lines 1-15). However, Willner et al. in view of Svedhem et al. and Bentley et al. fails to teach a coated metal surface on a solid support, wherein the antigens are bound to the same or different monolayers in patches on the solid support.

Duffy teaches an array system which can be used to elucidate interactions between molecules (p5, paragraph [0039]). The system comprises array of binding areas (patches) for immobilizing biomolecules and provides for high throughput, as many interactions may be tested in a single assay (p5, paragraphs [0040] and [0041]). Duffy further teaches that the interactions between molecules can be detected using QCM (p13, paragraph [0113]).

Therefore, it would have been obvious to one of ordinary skill in the art at the time of the invention to include an array of binding patches for immobilization of antigens of the Willner et al. in view of Svedhem et al. and Bentley et al. as taught by Duffy in order to perform high throughput analysis of many interactions, which may be tested in a single assay. The advantage of having the capacity to perform high throughput analysis of many interactions, which may be tested in a single assay, provides the motivation to include an array of binding patches for immobilization of

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antigens of the Willner et al. in view of Svedhem et al. and Bentley et al. with a reasonable expectation of success since Duffy teaches that the array system can be used with QCM detection methods to detect binding interaction on the array surface.

With respect to claim 4, Willner et al. teaches derivatized explosives, which include trinitrotoluene (TNT) and dinitrotoluene (DNT, Fig's 1A and 1B). The derivatized DNT includes an amine group, which form an amide linkage with ethylene glycol of OEG.

With respect to claim 5, Willner et al. teaches antigens are selected from cocaine and heroine (p5, lines 13-15).

Response to Arguments

15. Rejection of claims 1, 2, 6, and 7 under 35 U.S.C. 103(a) as being unpatentable over Willner et al. in view of Svedhem et al. and Bentley et al.

Applicant's arguments filed on October 30, 2006 have been fully considered but they are not persuasive in view of previously stated grounds of rejection.

Applicants' argument that there is no suggestion of any "capturing agent" other than cystamine (p10) in Willner et al. is not fond persuasive as Willner et al. teaches that any method intended for sensing the presence of explosive molecules or other types of low molecular weight molecules such as drugs should be highly sensitive and adapted for detecting a small amount of molecules (p5).

Applicant's argument regarding the use of the currently recited invention in displacement assays is not found persuasive as the device of Willner et al. can be also

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used in displacement assays, in which antigens are immobilized on the coated metal surface on a solid support (p24, lines 6-19 and Fig. 7).

In response to applicant's argument that there is no suggestion/motivation to combine the references, the examiner recognizes that obviousness can only be established by combining or modifying the teachings of the prior art to produce the claimed invention where there is some teaching, suggestion, or motivation to do so found either in the references themselves or in the knowledge generally available to one of ordinary skill in the art. See In re Fine, 837 F.2d 1071, 5 USPQ2d 1596 (Fed. Cir. 1988) and In re Jones, 958 F.2d 347, 21 USPQ2d 1941 (Fed. Cir. 1992). In this case, the advantage of having a structurally stable SAM, which has the characteristic of reducing nonspecific binding of proteins and other bioactive molecules provides the motivation to include the SAM of OEG-terminated alkanethiol amides of Svedhem et al. in the QCM biosensor of Willner et al. The advantage of using structurally stable SAM (p4495, Introduction, last paragraph), which has the characteristic of reducing nonspecific binding of proteins and other bioactive molecules (p4494, Introduction, first paragraph) is disclosed in Svedhem et al. Therefore, Applicants' argument that use of Examiner's own knowledge in stating that the advantage of having a structurally stable SAM, which has the characteristics of reducing nonspecific binding of proteins and other bioactive molecules as a motivation is not found persuasive in light of reasons set forth above.

16. Rejection of claims 3-5 under 35 U.S.C. 103(a) as being unpatentable over Willner et al. in view of Svedhem et al. and Bentley et al., and further in view of Duffy

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Applicant's arguments filed on October 30, 2006 have been fully considered but they are not persuasive in view of previously stated grounds of rejection and reasons set forth above.

17. Since prior art fulfills all the limitations currently recited in the claims, the invention as currently recited would read upon the prior art.

Conclusion

- 18. No claim is allowed.
- 19. **THIS ACTION IS MADE FINAL.** Applicant is reminded of the extension of time policy as set forth in 37 CFR 1.136(a).

A shortened statutory period for reply to this final action is set to expire THREE MONTHS from the mailing date of this action. In the event a first reply is filed within TWO MONTHS of the mailing date of this final action and the advisory action is not mailed until after the end of the THREE-MONTH shortened statutory period, then the shortened statutory period will expire on the date the advisory action is mailed, and any extension fee pursuant to 37 CFR 1.136(a) will be calculated from the mailing date of the advisory action. In no event, however, will the statutory period for reply expire later than SIX MONTHS from the mailing date of this final action.

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20. Any inquiry concerning this communication or earlier communications from the examiner should be directed to Unsu Jung whose telephone number is 571-272-8506. The examiner can normally be reached on M-F: 9-5.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Long Le can be reached on 571-272-0823. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free). If you would like assistance from a USPTO Customer Service Representative or access to the automated information system, call 800-786-9199 (IN USA OR CANADA) or 571-272-1000.

Unsu Jung, Ph.D. Patent Examiner Art Unit 1641

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